polycythemia vera

Polycythemia vera is a condition characterized by an increased number of red blood cells in the bloodstream. Affected individuals may also have excess white blood cells and blood clotting cells called platelets. These extra cells cause the blood to be thicker than normal. As a result, abnormal blood clots are more likely to form and block the flow of blood through arteries and veins. Individuals with polycythemia vera have an increased risk of deep vein thrombosis (DVT), a type of blood clot that occurs in the deep veins of the arms or legs. If a DVT travels through the bloodstream and lodges in the lungs, it can cause a life-threatening clot known as a pulmonary embolism (PE). Affected individuals also have an increased risk of heart attack and stroke caused by blood clots in the heart and brain.

Polycythemia vera typically develops in adulthood, around age 60, although in rare cases it occurs in children and young adults. This condition may not cause any symptoms in its early stages. Some people with polycythemia vera experience headaches, dizziness, ringing in the ears (tinnitus), impaired vision, or itchy skin. Affected individuals frequently have reddened skin because of the extra red blood cells. Other complications of polycythemia vera include an enlarged spleen (splenomegaly), stomach ulcers, gout (a form of arthritis caused by a buildup of uric acid in the joints), heart disease, and cancer of blood-forming cells (leukemia).

Frequency

The prevalence of polycythemia vera varies worldwide. The condition affects an estimated 44 to 57 per 100,000 individuals in the United States. For unknown reasons, men develop polycythemia vera more frequently than women.

Genetic Changes

Mutations in the *JAK2* and *TET2* genes are associated with polycythemia vera. Although it remains unclear exactly what initiates polycythemia vera, researchers believe that it begins when mutations occur in the DNA of a hematopoietic stem cell. These stem cells are located in the bone marrow and have the potential to develop into red blood cells, white blood cells, and platelets. *JAK2* gene mutations seem to be particularly important for the development of polycythemia vera, as nearly all affected individuals have a mutation in this gene. The *JAK2* gene provides instructions for making a protein that promotes the growth and division (proliferation) of cells. The JAK2 protein is especially important for controlling the production of blood cells from hematopoietic stem cells.

JAK2 gene mutations result in the production of a JAK2 protein that is constantly turned on (constitutively activated), which increases production of blood cells and prolongs their survival. With so many extra cells in the bloodstream, abnormal blood clots are more likely to form. Thicker blood also flows more slowly throughout the body, which prevents organs from receiving enough oxygen. Many of the signs and symptoms of polycythemia vera are related to a shortage of oxygen in body tissues.

The function of the *TET2* gene is unknown. Although mutations in the *TET2* gene have been found in approximately 16 percent of people with polycythemia vera, it is unclear what role these mutations play in the development of the condition.

Inheritance Pattern

Most cases of polycythemia vera are not inherited. This condition is associated with genetic changes that are somatic, which means they are acquired during a person's lifetime and are present only in certain cells.

In rare instances, polycythemia vera has been found to run in families. In some of these families, the risk of developing polycythemia vera appears to have an autosomal dominant pattern of inheritance. Autosomal dominant inheritance means that one copy of an altered gene in each cell is sufficient to increase the risk of developing polycythemia vera, although the cause of this condition in familial cases is unknown. In these families, people seem to inherit an increased risk of polycythemia vera, not the disease itself.

Other Names for This Condition

- Osler-Vaquez disease
- polycythemia ruba vera
- primary polycythemia
- PRV
- PV

Diagnosis & Management

Genetic Testing

 Genetic Testing Registry: Polycythemia vera https://www.ncbi.nlm.nih.gov/gtr/conditions/C0032463/

Other Diagnosis and Management Resources

- MedlinePlus Encyclopedia: Polycythemia Vera https://medlineplus.gov/ency/article/000589.htm
- MPN Research Foundation: Diagnosis http://www.mpnresearchfoundation.org/Polycythemia-Vera-28PV-29

General Information from MedlinePlus

- Diagnostic Tests
 https://medlineplus.gov/diagnostictests.html
- Drug Therapy https://medlineplus.gov/drugtherapy.html
- Genetic Counseling https://medlineplus.gov/geneticcounseling.html
- Palliative Care https://medlineplus.gov/palliativecare.html
- Surgery and Rehabilitation https://medlineplus.gov/surgeryandrehabilitation.html

Additional Information & Resources

MedlinePlus

- Encyclopedia: Polycythemia Vera https://medlineplus.gov/ency/article/000589.htm
- Health Topic: Blood Disorders https://medlineplus.gov/blooddisorders.html
- Health Topic: Bone Marrow Diseases https://medlineplus.gov/bonemarrowdiseases.html

Genetic and Rare Diseases Information Center

 Polycythemia vera https://rarediseases.info.nih.gov/diseases/7422/polycythemia-vera

Additional NIH Resources

- National Cancer Institute https://www.cancer.gov/types/myeloproliferative/patient/chronic-treatment-pdq#section/_225
- National Heart Lung and Blood Institute https://www.nhlbi.nih.gov/health/health-topics/topics/poly/

Educational Resources

- Disease InfoSearch: Polycythemia vera http://www.diseaseinfosearch.org/Polycythemia+vera/5826
- Leukemia and Lymphoma Society: Polycythemia Vera Facts http://www.lls.org/sites/default/files/file_assets/FS13_PolycythemiaVera_FactSheet.pdf

- MalaCards: polycythemia vera, somatic http://www.malacards.org/card/polycythemia_vera_somatic
- Merck Manual Home Health Handbook for Patients and Caregivers
 http://www.merckmanuals.com/home/blood-disorders/myeloproliferative-disorders/polycythemia-vera
- Merck ManualProfessional Version
 http://www.merckmanuals.com/professional/hematology-and-oncology/myeloproliferative-disorders/polycythemia-vera
- Orphanet: Polycythemia vera http://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=729

Patient Support and Advocacy Resources

- Leukemia and Lymphoma Society http://www.lls.org/
- MPN Research Foundation http://www.mpnresearchfoundation.org/
- National Organization for Rare Disorders (NORD)
 https://rarediseases.org/rare-diseases/polycythemia-vera/

ClinicalTrials.gov

ClinicalTrials.gov
 https://clinicaltrials.gov/ct2/results?cond=%22polycythemia+vera%22

Scientific Articles on PubMed

PubMed

https://www.ncbi.nlm.nih.gov/pubmed?term=%28Polycythemia+Vera%5BMAJR%5D%29+AND+%28polycythemia+vera%5BTI%5D%29+AND+review%5Bpt%5D+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1800+days%22%5Bdp%5D

OMIM

 POLYCYTHEMIA VERA http://omim.org/entry/263300

Sources for This Summary

- Cario H, McMullin MF, Pahl HL. Clinical and hematological presentation of children and adolescents with polycythemia vera. Ann Hematol. 2009 Aug;88(8):713-9. doi: 10.1007/s00277-009-0758-y. Epub 2009 May 26. Review.
 - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19468728
 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4135082/
- Delhommeau F, Jeziorowska D, Marzac C, Casadevall N. Molecular aspects of myeloproliferative neoplasms. Int J Hematol. 2010 Mar;91(2):165-73. doi: 10.1007/s12185-010-0530-z. Epub 2010 Feb 27. Review.
 - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20186505
- Gangat N, Strand J, Lasho TL, Finke CM, Knudson RA, Pardanani A, Li CY, Ketterling RP, Tefferi A. Cytogenetic studies at diagnosis in polycythemia vera: clinical and JAK2V617F allele burden correlates. Eur J Haematol. 2008 Mar;80(3):197-200. Epub 2007 Dec 7.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18081705
- Landolfi R, Nicolazzi MA, Porfidia A, Di Gennaro L. Polycythemia vera. Intern Emerg Med. 2010 Oct;5(5):375-84. doi: 10.1007/s11739-010-0369-6. Epub 2010 Mar 16. Review.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20237866
- Mehta J, Wang H, Iqbal SU, Mesa R. Epidemiology of myeloproliferative neoplasms in the United States. Leuk Lymphoma. 2014 Mar;55(3):595-600. doi: 10.3109/10428194.2013.813500. Epub 2013 Jul 29.
 - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/23768070
- Tefferi A. Novel mutations and their functional and clinical relevance in myeloproliferative neoplasms: JAK2, MPL, TET2, ASXL1, CBL, IDH and IKZF1. Leukemia. 2010 Jun;24(6):1128-38. doi: 10.1038/leu.2010.69. Epub 2010 Apr 29. Review.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20428194
 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3035972/
- Thiele J, Kvasnicka HM. The 2008 WHO diagnostic criteria for polycythemia vera, essential thrombocythemia, and primary myelofibrosis. Curr Hematol Malig Rep. 2009 Jan;4(1):33-40. doi: 10.1007/s11899-009-0005-6.

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